



## EU Declaration of Conformity

2625 Patton Road  
Roseville, MN 55113  
651-294-2200  
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This declaration of conformity is issued under the sole responsibility of the manufacturer. The device covered by the present declaration is in conformity with all regulations below, including compliance with related Essential Requirements.

### Object of the declaration:

<b>Product Name:</b>	<i>Powerlink 4 CE</i>
<b>Product Model Designator:</b>	Powerlink 4 CE
<b>Product Part Number(s):</b>	10010702
<b>Basic UDI-DI</b> <required for MDR>	00186648000449
<b>Control Indicator:</b>	2023W01 thru 2028W01
<b>Global Medical Device Nomenclature Code (GMDN) and Description</b>	
<b>Product Options/Accessories:</b>	N/A

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The object of the declaration described above is in conformity with the following regulations:

<b>EU Regulation</b>	<b>Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices</b>
<b>Device Classification:</b>	Class I based on Annex VIII and Rule 13
<b>Conformity Assessment Path</b>	<i>Class I device – Self Certify – per MDR 2017/745 Annex II and III</i>
<b>Name/Address/ID of Notified Body:</b>	<i>Not Applicable – Class I device</i>
<b>Standards and Common Specifications</b>	<p>The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below.</p> <p><i>BS EN ISO 14971:2019+A11:2021 – Medical Devices – Application of risk management to medical devices</i>  <i>BS EN ISO 13485:2016+A11:2021 – Medical Devices – Quality management systems – Requirements for regulatory purposes</i>  <i>EN 60601-1-1:1998:A1: 1991 + A2:1995</i>  <i>EN 60730-1:2016 : EN 60730-2-7:2010</i></p>


<b>EU Regulation</b>	<b>Radio Equipment Directive (2014/53/EU)</b>
<b>Standards and Common Specifications</b>	<p>The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below.</p> <p><i>ETSI EN 300 440 V2.1.1 (2017-03)</i>  <i>ETSI EN 301 489-1 V2.2.3 (2019-11)</i>  <i>ETSI EN 301 489-3 V2.1.1 (2017-03)</i></p>

<b>EU Regulation</b>	<b>Restriction of Hazardous Substances in Electrical and Electronic Equipment Directive 2011/65/EU</b>
<b>Standards and Common Specifications</b>	<p>The products listed above have been tested in a typical configuration as described in the Manufacturer's accompanying documentation and are compliant with the product standards listed below.</p> <p><i>Supply and Component Controls - CPSC-CH-E1002-08.1 and/or CPSC-CH-E1001-08.1 and CPSC-CH-C1001-09.3</i></p>

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**Additional information:**

<b>EU Authorized Representative:</b>	<i>EUCEREP B.V. Roald Dahllaan 33 5629MC – Eindhoven The Netherlands</i>
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Signature (signed for and on behalf of AbleNet): 	Date of Issue: <DD Month YYYY> 13 January 2023
Printed Name: Joe Volp	Place of Issue: AbleNet Inc – Roseville, MN
Title: Director of Marketing	Document Number: DoC_10010702_Powerlink 4 CE_011623